### **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

### 1-20. (canceled)

- 21. (previously presented) The amyloid-targeting imaging agent of claim 31, wherein A<sub>lab</sub> includes a radionuclide selected from <sup>99m</sup>Tc, <sup>99</sup>Tc, <sup>64</sup>Cu, <sup>67</sup>Cu, <sup>97</sup>Ru, <sup>109</sup>Pd, <sup>186</sup>Re, <sup>188</sup>Re, <sup>111</sup>In, <sup>113m</sup>In, <sup>153</sup>Gd, <sup>90</sup>Y, <sup>153</sup>Sm, <sup>166</sup>Ho, <sup>198</sup>Au, <sup>199</sup>Au, <sup>90</sup>Sr, <sup>89</sup>Sr, <sup>105</sup>Rh, <sup>201</sup>Tl, <sup>51</sup>Cr, <sup>67</sup>Ga, <sup>57</sup>Co, <sup>60</sup>Co, <sup>123</sup>I, <sup>125</sup>I, <sup>131</sup>I or <sup>18</sup>F.
- 22. (previously presented) The amyloid-targeting imaging agent of claim 31, wherein A<sub>lab</sub> includes a radionuc lide selected from the group consisting of Tc and Re.
- 23. (previously presented) The amyloid-targeting imaging agent of claim 31, wherein A<sub>lab</sub> is a metal chelate of a radioactive or paramagnetic metal ion.
- 24. (withdrawn) The amyloid-targeting imaging agent of claim 31, wherein A<sub>lab</sub> comprises a chelating ligand of the formula

where  $R^{10}$  is a linear or branched, saturated or unsaturated  $C_{1.4}$  alkylene group interrupted by one or two heteroatoms;  $R^{11}$  is H or  $R^{10}$ , or  $R^{10}$  and  $R^{11}$  taken together, form a 5- to 8-membered saturated or unsaturated heterocyclic ring optionally substituted with one or more of halogen, hydroxyl, amino, carboxyl, oxo,  $C_{1-4}$  alkyl, aryl, or C(O)R groups;  $R^3$ ,  $R^4$ ,  $R^5$  and  $R^6$  are independently H, carboxyl,  $C_{1-4}$  alkyl, an alpha carbon side chain of a D- or L-amino acid other than proline, or C(O)R;  $R^7$  and  $R^8$  are independently H,

carboxyl, amino,  $C_{1-4}$  alkyl,  $C_{1-4}$  alkyl;  $R^9$  is H or a sulfur protecting group; and L is hydroxyl, alkoxy, an amino acid residue, or a linking group.

# 25-30. (canceled)

# 31. (currently amended) An amyloid-targeting imaging agent of the formula

$$A_{t} - A_{lnk} - A_{lab}$$
 (I)

where z is 0 or 1; At is an amyloid targeting moiety selected from the group consisting of

NaO<sub>3</sub>SCH<sub>2</sub>(CH<sub>2</sub>)<sub>4</sub>CH<sub>2</sub>SO<sub>3</sub>Na, NaO<sub>3</sub>SNHCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OSO<sub>3</sub>Na, NaO<sub>3</sub>SNHCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OSO<sub>3</sub>Na,

$$H_3C \longrightarrow SO_3Na$$

and pharmaceutically acceptable salts thereof,

A<sub>lnk</sub> is a linker moiety; and A<sub>lab</sub> is a labeling moiety.

### 32-42. (canceled)

- 43. (previously presented) A kit for preparing a radiopharmaceutical preparation, said kit comprising:
  - an amyloid-targeting imaging agent of claim 31;
  - a reducing agent;
  - a buffering agent;
  - a transchelating agent, and

instructions for the preparation and use of the radiopharmaceutical in the imaging of amyloid or an amyloid-related condition.

### 44-50. (canceled)

- 51. (withdrawn) A method of diagnostic medical imaging of an amyloid-associated disease comprising the steps of administering to a patient a pharmaceutical composition according to claim 31 and then imaging said patient.
- 52. (withdrawn) The method of diagnostic medical imaging according to claim 51 wherein A<sub>lab</sub> of said pharmaceutical composition is a radiopharmaceutical.

53. (withdrawn) The method of diagnostic medical imaging according to claim 51 wherein A<sub>lab</sub> of said pharmaceutical composition is a metal chelate.

- 54. (withdrawn) The method of diagnostic medical imaging according to claim 53 wherein said metal chelate is gadolinium-DTPA, gadolinium-DOTA, or gadolinium-DO3A.
- 55. (withdrawn) The method of diagnostic medical imaging according to claim 53 wherein said metal chelate is a chelate of <sup>99m</sup>Tc or <sup>111</sup>In.
- 56. (withdrawn) The method of diagnostic medical imaging according to claim 51 wherein said imaging step is ultrasound imaging.
- 57. (withdrawn) The method of claim 105, wherein said imaging step is radionuclide imaging.
- 58. (withdrawn) The method of claim 57, wherein said imaging step is SPECT imaging.
- 59. (withdrawn) The method of claim 105, wherein said imaging step is magnetic resonance imaging.
- 60. (withdrawn) The method of claim 105, wherein said imaging step is ultrasound imaging.
- 61. (withdrawn) The method of claim 105, wherein said imaging step is X-ray imaging.
- 62. (withdrawn) The method of claim 105, wherein said imaging step is fluorescence imaging.
- 63-94. (canceled)
- 95. (withdrawn) A method for diagnostic medical imaging of an amyloid-associated disease in a patient, comprising administering to a patient a pharmaceutical composition comprising an amyloid-targeting imaging agent of claim 31, and imaging the amyloid-targeting imaging agent in said patient.
- 96. (withdrawn) The method of claim 95, wherein A<sub>lab</sub> of said pharmaceutical composition is a radiopharmaceutical.

97. (withdrawn) The method of claim 95, wherein A<sub>lab</sub> of said pharmaceutical composition is a metal chelate.

- 98. (withdrawn) The method of claim 95, wherein A<sub>lab</sub> of said pharmaceutical composition is a metal chelate and said imaging step is magnetic resonance imaging or radionuclide imaging.
- 99. (withdrawn) The method of claim 97, wherein said metal chelate is gadolinium-DTPA, gadolinium-DOTA, or gadolinium-DO3A.
- 100. (withdrawn) The method of claim 97, wherein said metal chelate is a chelate of <sup>99m</sup>Tc or <sup>111</sup>In.
- 101. (withdrawn) The method of claim 95, wherein said imaging step is ultrasound imaging.
- 102. (canceled)
- 103. (withdrawn) A method for diagnosing an amyloid-related condition in a patient, comprising administering an amyloid-targeting imaging agent according to claim 31 to a patient, and imaging said amyloid-targeting imaging agent in said patient to determine the presence of amyloid in said patient, such that the presence or absence of an amyloid-related condition in said patient is determined.
- 104. (withdrawn) The method of claim 103, wherein said amyloid-related condition is selected from the group consisting of Creutzfeld-Jakob Disease (CJD), Kuru, transmissible cerebral amyloidosis-amyloidosis, (also known as transmissible virus dementias[[)]], familial CJD, scrapie, transmissible mink encephalopathy, bovine spongiform encephalopathy (BSE), inflammation-associated amyloid, type II diabetes, primary amyloidosis, feline spongiform encephalopathy, non-transmissible cerebral amyloidosis (e.g., Alzheimer's disease), prion-mediated diseases, dialysis-related amyloidosis, light chain-related amyloidosis, cerebral amyloid angiopathy, and Alzheimer's disease.
- 105. (withdrawn) A method for imaging amyloid deposition in a patient, comprising administering an amyloid-targeting imaging agent according to claim 31 to a patient, and

imaging said amyloid-targeting imaging agent in said patient to determine the presence of amyloid in said patient.

106-131. (canceled)

132. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

$$NaO_3SCH_2(CH_2)_4CH_2SO_3Na$$

or a pharmaceutically acceptable salt thereof.

133. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

# NaO3SNHCH2CH2CH2OSO3Na

## NaO<sub>3</sub>SNHCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OSO<sub>3</sub>Na

or a pharmaceutically acceptable salt thereof.

134. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

or a pharmaceutically acceptable salt thereof.

135. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

# or a pharmaceutically acceptable salt thereof.

136. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

# or a pharmaceutically acceptable salt thereof.

137. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

### or a pharmaceutically acceptable salt thereof.

138. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

# or a pharmaceutically acceptable salt thereof.

139. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

$$N$$
  $SO_3H$ 

### or a pharmaceutically acceptable salt thereof.

140. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

# or a pharmaceutically acceptable salt thereof.

141. (currently amended) The amyloid-targeting imaging agent of claim 31, wherein the amyloid targeting moiety is of the formula

$$Cl^{-}$$
 $CO_{2}Me$ 

# or a pharmaceutically acceptable salt thereof.